# INFORMATION DISCLOSURE STATEMENT BY APPLICANT ( Not for submission under 37 CFR 1.99)

Application Number		09560203	
Filing Date		2000-04-28	
First Named Inventor	Erisman		
Art Unit		3693	
Examiner Name	Danie	niel Felten	
Attornou Docket Number		TEE 2000-1	

#### CERTIFICATION STATEMENT

Please see 37	CFR 1	.97 and	1.98 to make the	appropriate	selection(s):
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That each item of information contained in the information disclosure statement was first cited in any communication from a foreign patient office in a counterpart foreign application not more than three months prior to the filing of the information disclosure statement. See 37 CFF 1.97(e)(1).

## OR

That no item of information contained in the information disclosure statement was cited in a communication from a foreign patent office in a counterpart foreign application, and, to the knowledge of the person signing the certification after making reasonable inquiry, no item of information contained in the information disclosure statement was known to any individual designated in 37 CFR 1.59(c) more than three months prior to the filing of the information disclosure statement. See 37 CFR 1.57(e)(2).

- See attached certification statement.
- Fee set forth in 37 CFR 1.17 (p) has been submitted herewith.

IOHN N. GROSS

☐ None

Name/Print

#### SIGNATURE

A signature of the applicant or representative is required in accordance with CFR 1.33, 10.18. Please see CFR 1.4(d) for the form of the signature.

form of the signature.						
Signature	/JOHN N. GROSS/	Date (YYYY-MM-DD)	2010-06-11			

Registration Number

34175

This collection of information is required by 3T CFR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is for lie and by the USPTO to process) an application. Confidentiality is governed by \$5 U.S. C. 12 and 37 CFR 1.14. This collection is estimated to take 1 hour to complete, including gathering, preparing and submitting the completed application from to the USPTO. There will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. operatment of Comments o

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The Privacy Act of 1974 (P. L. 95-79) requires that you be given certain information in connection with your submission of the attached form related to a patient application or patient. Accordingly, pursuant to the requirements of the Act, please be advised that: (1) the general authority for the collection of this information is 35 U.S.C. (2)(2)(2) furnishing of the information solicited to isolutionary, and (3) the principal purpose for which the information is used by the U.S. Patient and Trademan KORICs is to information, the U.S. Patient and Trademan KORICs may not be able to process and/or examine your submission, which may result in formation of proceedings or abandonment of the application or experigation of the patient.

The information provided by you in this form will be subject to the following routine uses:

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- A record related to an International Application filed under the Patent Cooperation Treaty in this system of records
  may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant
  to the Patent Cooperation Treaty.
- A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
- 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or hisher designed, cuting an inspection of records conducted by GSA is part of that apency's responsibility to recommend improvements in records management practices and programs, under authority of 4 U.S.C. 2904 and 2906. Such disclosure shall be made in accordance with the GSA regulations governing inspection of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make determinations abavit individuals.
- A record from this system of records may be disclosed, as a routine use, to the public after either publication of
  the application pursuant to 35 U.S. C. 12(b) or issuance of a patent pursuant to 35 U.S. C. 157. Twither, a record
  may be disclosed, subject to the imitiations of 37 CFR 1.14, as a routine use, to the public if the record was filed in
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